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Docket No.: 19338CDCPA2 Serial No.: 08/554,424

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## IN THE CLAIMS

1-23 (Previously cancelled or withdrawn)

FROM-BEUSSE BROWNLEE ET AL

- 24. (Currently Amended) A method of identifying a ligand that modulates activity of a Drosophila membrane voltage-activated sodium channel, which comprises:
  - expressing an isolated Drosophila voltage-activated sodium channel para, (a) and expressing an isolated Drosophila voltage-activated putative beta subunit, tipE, in a first Xenopus oocyte host cell, wherein said expressing of para and said expressing of tipE occur after coinjection of para and tipE RNA, wherein said para RNA is encoded by the DNA molecule as set forth in SEQ ID NO: 7, and wherein the host cell resultingly expresses a voltage-activated sodium current that is tetrodotoxin sensitive;
  - contacting the first host cell with said ligand; (b)
  - measuring the resulting voltage-activated current; and (c)
  - comparing the voltage-activated current measured according to step (c) (d) with voltage-activated current measured in a second, control Xenopus oocyte host cell prepared according to step (a) and not treated with said ligand.
- 25. (Currently Amended) A method of identifying a ligand that modulates activity of a Drosophila membrane voltage-activated sodium channel, which comprises:
- (a) co-expressing an isolated Drosophila voltage-activated sodium channel para and an isolated Drosophila voltage-activated putative beta subunit, tipE, in a host cell from a multicellular organism, wherein said co-expressing of pura and tipE occurs after an isolated DNA molecule encoding para and an isolated DNA molecule encoding upE are introduced into said host cell, wherein said isolated DNA molecule which encodes para is as set forth in SEQ ID NO: 7, and wherein the host cell resultingly expresses a voltage-activated sodium current that is tetrodotoxin sensitive;
- (b) contacting the first host cell with said ligand;
- (c) measuring the resulting voltage-activated current; and

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- (d) comparing the voltage-activated current measured according to step (c) with voltage-activated current measured in a second, control Xenopus oocyte host cell prepared according to step (a) and not treated with said ligand.
- 26. (Currently Amended) A method of identifying a ligand that modulates <u>activity of a Drosophila</u> membrane voltage-activated sodium channel, which comprises:
- (a) expressing an isolated *Drosophilu* voltage-activated sodium channel *para*, and expressing an isolated *Drosophilu* voltage activated putative beta subunit *tipE*, in a host cell selected from the group consisting of *Xenopus* oocytes and a cell from a multicellular organism, wherein an isolated DNA molecule which expresses *para* comprises a DNA sequence as set forth in SEQ ID NO: 7, and wherein the host cell resultingly expresses a voltage-activated sodium current that is tetrodotoxin sensitive;
- (b) contacting the first host cell with said ligand;
- (c) measuring the resulting voltage-activated current;
- (d) comparing the voltage-activated current measured according to step (c) with voltageactivated current measured in a second, control *Xenopus* oocyte host cell prepared according to step (a) and not treated with said ligand; and
- (e) comparing the voltage-activated current measured according to step (c) with voltage-activated current produced prior to contacting the host cell with the ligand.
- 27. (Previously Added) The method of claim 24, additionally comprising comparing the voltage-activated current measured according to step (c) with voltage-activated current measured upon contacting said ligand with a third control host cell in which said para and said tipE are not co-expressed.
- 28. (Previously Added) The method of claim 25, additionally comprising comparing the voltage-activated current measured according to step (c) with voltage-activated current measured upon contacting said ligand with a third control host cell in which said para and said tipE are not co-expressed.

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29. (Previously Added) The method of claim 26, additionally comprising comparing the voltage-activated current measured according to step (c) with voltage-activated current measured upon contacting said ligand with a third control host cell in which said para and said tipE are not co-expressed.